AMENDMENTS TO THE SPECIFICATION

Please amend the paragraph bridging pages 14-15 as follows:

The dose of the preparation of the invention may be conventionally selected such that it causes a satisfactory therapeutic response. In general, the dose may be usually selected from the range of about 0.01 to about **[[**10,00**]]** 10,000 μg, and preferably the range of about 0.1 to about 1000 μg per one adult patient. The preparation can be embedded or infused in the lesion or a peripheral site thereof. Furthermore, when the effect is insufficient with a single dose, additional dose may be administered several times.